McNeil, M.D.; Walter J. McNerney, M.H.A.; Edward B. Perrin, Ph.D; Louis F. Rossiter, Ph.D.; Albert L. Siu, M.D.; and Ellen B. White, M.B.A.

There also are Federal ex-officio members. These members are:
Administrator, Substance Abuse and Mental Health Services Administration; Director, National Institutes of Health; Director, Centers for Disease Control and Prevention; Administrator, Health Care Financing Administration; Commissioner, Food and Drug Administration; Assistant Secretary of Defense (Health Affairs); and Chief Medical Director, Department of Veterans Affairs.

II. Agenda

On Thursday, May 30, 1996, the meeting will begin at 9:00 a.m. with the call to order by the Council Chairman. The Administrator, AHCPR, will update the status of current Agency issues and program initiatives. The Council will then discuss issues concerning the guideline program, public/private sector

collaboration, and improvements to the large grant program. The Council will recess at 3:45 p.m. The Council will begin the closed portion of the meeting to review grant applications from 4:00 p.m. to 5:00 p.m.

The meeting will adjourn at 5:00 p.m. Agenda items are subject to change as priorities dictate.

Dated: May 8, 1996. Clifton R. Gaus, Administrator. [FR Doc. 96–12066 Filed 5–13–96; 8:45 am] BILLING CODE 4160–90–M

Administration for Children and Families

Agency Recordkeeping/Reporting Requirements Under Emergency Review by the Office of Management and Budget (OMB)

Title: Request for Emergency OMB Approval of Information Collection Under the Paperwork Reduction Act. OMB No.: New.

Description: The State and Tribal JOBS plans are statutorily mandated and serve as the agreement between the State or the Tribal grantee and the Federal government for how JOBS programs will operate. The State/Tribal plans provide assurances that the JOBS program will be administered and operated in conformity with titles IV-A and IV-F of the Social Security Act, pertinent Federal regulations, and other applicable instructions or guidelines issued by ACF. This new State and Tribal JOBS plan section is being added in response to the President's recent directive requiring States to address the needs of teen parents so that they stay in school and become self-sufficient.

Respondents: State, Local or Tribal Govt.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total bur- den hours
Tribal JOBSState JOBS	76 54	1 1	5 5	380 270

Estimated Total Annual Burden Hours: 650.

Additional Information

ACF is requesting that OMB grant a 90 day approval for this information collection under procedures for emergency processing by May 9, 1996. A copy of this information collection, with applicable supporting documentation, may be obtained by calling the Administration for Children and Families, Reports Clearance Officer, Roberta Katson at (202) 401–5756.

Comments and questions about the information collection described above should be directed to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ACF, Office of Management and Budget, Paperwork Reduction Project, 725 17th Street NW., Washington, DC 20503, (202) 395–7316.

Roberta Katson,

Director, Office of Information Resource

Management Services.

[FR Doc. 96–11946 Filed 5–13–96; 8:45 am]

BILLING CODE 4184–01–M

Dated: May 8, 1996.

Food and Drug Administration

FDA Form 3439, Interim Form for Application to Market a New Drug, Biologic, or Antibiotic Drug for Human Use: Availability

AGENCY: Food and Drug Administration, HHS

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of FDA Form 3439 entitled, "Application to Market a New Drug, Biologic, or Antibiotic Drug for Human Use." This form is intended for use by applicants for licenses for specified biotechnology and specified synthetic biological products. FDA Form 3439 has received interim approval from the Office of Management and Budget (OMB) for use by applicants pending the availability of a harmonized form for use by applicants requesting approval of drugs, biological products, and antibiotics. The development of the harmonized form and this action are part of FDA's continuing effort to achieve the objectives of the President's "Reinventing Government" initiatives.

EFFECTIVE DATE: The FDA Form 3439 may be used May 24, 1996.

ADDRESSES: Submit written requests for single copies of FDA Form 3439 to Division of Congressional and Public Affairs (HFM-44), Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852-1448. Send one self-addressed adhesive label to assist that office in processing your requests. The form may also be obtained by FAX by calling the CBER Voice Information System at 1–800-835-4709. FDA Form 3439 is available for public examination in the Dockets Management Branch (HFA-305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1-23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Annette A. Ragosta, Center for Biologics Evaluation and Research (HFM–630), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–594–3074.

SUPPLEMENTARY INFORMATION: FDA is making available FDA Form 3439, Application to Market a New Drug,